Enrotron Flavour 50 mg Tablets for dogs

Authorised

- Enrofloxacin

Product identification

Medicine name:
Enrotron Flavour 50 mg Tablets for dogs

Active substance:
Enrofloxacin

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Enrofloxacin
50.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Withdrawal period by route of administration:
Oral use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Package description:
(ID3) 30 Tablet: unspecified outer container with 1 Blister (Aluminium; Aluminium) with 30 Tablet
(ID9) 50 Tablet: unspecified outer container with 1 Blister (Aluminium; PolyVinyl Chloride) with 50 Tablet
(ID6) 10 Tablet: unspecified outer container with 1 Blister (Aluminium; PolyVinyl Chloride) with 10 Tablet
(ID2) 20 Tablet: unspecified outer container with 1 Blister (Aluminium; Aluminium) with 20 Tablet
(ID5) 100 Tablet: unspecified outer container with 1 Blister (Aluminium; Aluminium) with 100 Tablet
(ID7) 20 Tablet: unspecified outer container with 1 Blister (Aluminium; PolyVinyl Chloride) with 20 Tablet
(ID8) 30 Tablet: unspecified outer container with 1 Blister (Aluminium; PolyVinyl Chloride) with 30 Tablet
(ID4) 50 Tablet: unspecified outer container with 1 Blister (Aluminium; Aluminium) with 50 Tablet
(ID1) 10 Tablet: unspecified outer container with 1 Blister (Aluminium; Aluminium) with 10 Tablet
(ID10) 100 Tablet: unspecified outer container with 1 Blister (Aluminium; PolyVinyl Chloride) with 100 Tablet

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Animedica GmbH

Marketing authorisation date:
24/11/2010

Manufacturing sites for batch release:
Animedica GmbH

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
VM 24745/4012

Date of authorisation status change:
24/11/2010

Reference member state:
Germany

Procedure number:
DE/V/0137/001

Concerned member states:
Austria Belgium Denmark Finland Greece Hungary Iceland Ireland Luxembourg Netherlands Poland Slovenia United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

Source URL: *https://medicines.health.europa.eu/veterinary/600000064846*